AMENDMENTS TO THE CLAIMS

- (Currently amended) A process for the preparation of a composite containing a
 drug dispersed in an organic carrier, wherein the drug is massively dispersed (in bulk) within the
 particles of said organic carrier and it is present in amorphous form in a quantity greater than or
 equal to 50%; comprising the following steps:
- a) forming a mixture of mixing a drug with an and a particulate organic carrier selected from the group consisting of water-soluble complexing agents ehesen from evelodextrins and maltodextrins, water-insoluble cross-linked polymers, and mixtures thereof;
- b) irradiating applying an oscillating electromagnetic field to the mixture obtained in a), with microwaves, wherein the oscillating electromagnetic field is microwave irradiation microwave power is modulated so that to increase the temperature of the mixture increases until it reaches a value higher to a temperature greater than the melting temperature of the drug and it is then maintained constant at the temperature greater than the melting temperature of the drug said value for at least 5 minutes to provide a composite containing the drug dispersed within the particulate organic carrier deposited both on the surfaces and inside the organic carrier particles, wherein the drug is present in the composite in amorphous form in a quantity greater than or equal to 50% by weight based on the total amount of the drug.
- (Currently amended) Process according to The process of claim 1, wherein in
 step a) a wet mixture is formed by mixing a drug and a particulate organic carrier further
 comprises adding a solvent to provide a wet mixture.
- (Currently amended) Process-according to <u>The process of claim 2</u>, wherein said solvent is water

LAW OFFICES OF CHRISTENSEN CYCONNOR JOHNSON KINDNESS*** 1420 Fifth Avenue Suite 2800 Seattle, Washington 98101 20.66.82.8100 4. (Currently amended) The process aecording to of claim 3, in which wherein said wet mixture is formed by adding water to the earrier-drug composite drug and particulate organic carrier in a quantity comprised of between 0.1 ml/g and 5 ml/g with respect to the dry-mixture of the composite based on the weight of drug and particulate organic carrier.

5. (Currently amended) The process according to of claim 2, in which wherein the oscillating electromagnetic field is applied to the mixture at a the pressure at which the irradiation is carried out is comprised of between 1 and 20 bar.

6. (Currently amended) A process according to <u>The process of claim 1</u>, wherein step b) the oscillating electromagnetic field is applied to the mixture is carried out in a container constituted of comprising a dielectric material having coupling capacity with [[the]] microwaves.

(Currently amended) The process aecording to of claim 6, wherein said dielectric
material is polytetrafluoroethylene loaded with graphite.

8. (Currently amended) The process according to of claim 1, in which wherein the microwave irradiation with microwaves is carried out with power in a power the range comprised of between 100 W and 5000 W, for an overall a time up to 120 minutes.

9. (Currently amended) A-process-according to The process of claim 1, wherein said cross-linked polymer is selected from the group consisting of cross-linked polyvinylpyrrolidone, cross-linked sodium carboxymethylcellulose, cross-linked starch, cross-linked dextran, cross-linked polystyrene and cross-linked β -cyclodextrin.

 (Currently amended) A process according to <u>The process of</u> claim 1, wherein said drug is a drug sparingly soluble in water.

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11-20. (Canceled)

 (New) The process of claim 1, wherein said water-soluble complexing agents are selected from the group consisting of cyclodextrins, maltodextrins, and mixtures thereof.